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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/527,101	12/27/2005	Sarah C. Bodary-Winter	P1978R1	3802	
9157 7590 08/08/2007 GENENTECH, INC. 1 DNA WAY SOUTH SAN FRANCISCO, CA 94080			EXAMINER		
			BASKAR, PADMAVATHI		
			ART UNIT	PAPER NUMBER	
			1645		
			[		
			MAIL DATE	DELIVERY MODE	
			08/08/2007	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)					
	10/527,101	BODARY-WINTER ET AL.					
Office Action Summary	Examiner	Art Unit					
•	Padmavathi v. Baskar	1645					
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	orrespondence address					
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA  - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication.  - If NO period for reply is specified above, the maximum statutory period was prepared to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tim vill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONEI	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).					
Status							
1) Responsive to communication(s) filed on							
2a) This action is <b>FINAL</b> 2b) This	action is non-final.						
3) Since this application is in condition for allowar	3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is						
closed in accordance with the practice under E	x parte Quayle, 1935 C.D. 11, 45	53 O.G. 213.					
Disposition of Claims							
4) ⊠ Claim(s) 1-26 is/are pending in the application. 4a) Of the above claim(s) is/are withdray 5) □ Claim(s) is/are allowed. 6) □ Claim(s) is/are rejected. 7) □ Claim(s) is/are objected to. 8) ⊠ Claim(s) 1-26 are subject to restriction and/or expressions.	vn from consideration.						
Application Papers							
9) The specification is objected to by the Examine	r	·					
10) The drawing(s) filed on is/are: a) acce		Examiner.					
Applicant may not request that any objection to the	drawing(s) be held in abeyance. See	e 37 CFR 1.85(a).					
Replacement drawing sheet(s) including the correcting 11) The oath or declaration is objected to by the Ex	- · · · · · · · · · · · · · · · · · · ·	`					
Priority under 35 U.S.C. § 119							
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of:  1. Certified copies of the priority documents 2. Certified copies of the priority documents 3. Copies of the certified copies of the prior application from the International Bureau * See the attached detailed Office action for a list of	s have been received. s have been received in Application ity documents have been receive (PCT Rule 17.2(a)).	on No ed in this National Stage					
220 mg sites as a samual since as an ion a list.	2 22 34 35 plas flot (300)	<del>-</del>					
Attachment(s)							
1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:	nte					

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1. Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group I, Claims 1-9 (in part) are drawn to an isolated polynucleotide, recombinant expression vector, host cell, and a process for producing polypeptide

Further restriction to one SEQ.ID.NO required (see paragraph #4).

Group II, Claims 10-12 and 15-17(in part) drawn to polypeptide and a composition comprising said polypeptide.

Further restriction to one SEQ.ID.NO required (see paragraph # 4).

Group III Claims 13-14 and 15 –17 (in part) drawn to an antibody and a composition comprising said polypeptide agonist or antagonist or antibody

Further restriction to one SEQ.ID.NO required (see paragraph # 4).

Group IV Claims 18 (in part) drawn to an article manufacture Further restriction to one SEQ.ID.NO required (see paragraph # 4).

Group V Claim 19 (in part) drawn to a method for treating psoriasis using PRO polypeptide or antagonist or antibody

Further restriction to one SEQ.ID.NO required (see paragraph # 4).

Groups VI Claim 20 (in part) drawn to a method for determining the presence of PRO polypeptide in a sample using anti PRO antibody

Further restriction to one anti PRO- antibody required (see paragraph # 4).

Group VII Claim 21 (in part) drawn to a method for diagnosing psoriasis using gene encoding PRO polypeptide

Further restriction to one PRO polypeptide required (see paragraph # 4).

Group VIII Claim 22 (in part) drawn to a method for diagnosing psoriasis using anti pro antibody Further restriction to one anti PRO- antibody required (see paragraph # 4).

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Group IX Claim 23 (in part) drawn to a method of identifying a compound that inhibits the activity of PRO polypeptide

Further restriction to one PRO-polypeptide required (see paragraph # 4).

Group X Claims 24-25 (in part) drawn to a method of identifying a compound that inhibits the expression of gene enclosing PRO-polypeptide

Further restriction to one PRO- polypeptide required (see paragraph # 4).

Group XI Claim 26 (in part) drawn to a method of identifying a compound that mimics the activity of PRO polypeptide

Further restriction to one PRO- polypeptide required (see paragraph # 4).

The inventions listed as Groups 1-XI (in Part) do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

The technical feature of linking Groups I-XI appears to be that they all relate to polynucleotide, polypeptide and antibody.

The technical feature of linking groups appears to be that they are all related to peptides, nucleic acids and antibodies and methods of using peptides, nucleic acids and antibodies.

However, Baker KP, et al WO200168848-A2disclose an isolated nucleic acid enclosing SEQ.ID.NO:2 (see the sequence analysis below)

## Alignment Scores:

Pred. No.:

1.2e-85 Length: 732

Score:

895.00

Matches:

160

Percent Similarity:

100.0%

Conservative: 0

Best Local Similarity: 100.0%

Mismatches: 0

Query Match:

100.0%

Indels:

0

DB:

4

Gaps:

US-10-527-101A-2 (1-160) x AAS46213 (1-732)

 ${\tt 1~MetIleAsnProGluLeuArgAspGlyArgAlaAspGlyPheIleHisArgIleValPro~20}\\$ Qу Вb 19 ATGATTAATCCAGAGCTGCGGGATGGCAGAGCTGATGGCTTCATACATCGGATAGTTCCC 78 21 LysLeuIleGlnAsnTrpLysIleGlyLeuMetCysPheLeuSerIleIleIleThrThr 40 Qy 

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Db	79	AAGTTGATACAAAACTGGAAGATTGGCCTTATGTGCTTCCTGAGTATTATTATTACTACA	138
Qу	41	ValCysIleIleMetIleAlaThrTrpSerLysHisAlaLysProValAlaCysSerGly	60
Db	139	GTTTGCATTATTATGATAGCCACATGGTCCAAGCATGCTAAACCTGTGGCATGTTCAGGG	198
Qу	61	AspTrpLeuGlyValArgAspLysCysPheTyrPheSerAspAspThrArgAsnTrpThr	80
Db	199	GACTGGCTTGGAGTGAGAGATAAGTGTTTCTATTTTTCTGATGATACCAGAAATTGGACA	258
Qу	81	${\tt AlaSerLysIlePheCysSerLeuGlnLysAlaGluLeuAlaGlnIleAspThrGlnGluing}$	100
Db	259	GCCAGTAAAATATTTTGTAGTTTGCAGAAAGCAGAACTTGCTCAGATTGATACACAAGAA	318
Qу	101	${\tt AspMetGluPheLeuLysArgTyrAlaGlyThrAspMetHisTrpIleGlyLeuSerArg}$	120
		111111111111111111111111111111111111	
Db	319	GACATGGAATTTTTGAAGAGGTACGCAGGAACTGATATGCACTGGATTGGACTAAGCAGG	378
Qу	121	LysGlnGlyAspSerTrpLysTrpThrAsnGlyThrThrPheAsnGlyTrpProSerAsn	140
DЬ	379	AAACAAGGAGATTCTTGGAAATGGACAAATGGCACCACATTCAATGGTTGGCCATCAAAC	438
Qу	141	SerLysTrpSerCysAsnTrpSerLeuArgGlnTrpLeuLeuLeuGlyProLeuArg	160
Db	439	TCCAAATGGTCTTGCAACTGGAGCCTCCGACAATGGCTTCTTCTGCTGGGACCCCTTAGA	498

and thus read on group I. Therefore, the technical feature of linking groups I-XI does not constitute a special technical feature as defined by PCT Rule 13.2, as it does not define a contribution over the prior art and hence unity of invention is lacking.

3. The special technical feature of Group I considered to be nucleic acid encoding polypeptide that shares no common structure, property and function with Group II since peptides contain amino acids and do not share the same or a corresponding technical feature with Group II nucleic acids.

The special technical feature of Group III considered to be antibody that shares no common structure, property and function from Inventions I and II since it has an inherent affinity, avidity, and specificity that DNA or a simple protein is not capable of expressing and do not require each other for their practice.

The technical feature linking groups IV-XI is considered to be methods utilizing products that share no common structure, property and function so as to form a single general inventive concept under Rule 13.1. Hence, unity is lacking among groups.

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4. For each group of inventions I-XI above, restriction to one of the following SEQ.ID.NO or PRO polypeptide or PRO antibody is also required under PCT Rule 13.1 because, under PCT Rule 13.2. Therefore, election is required of one of inventions I-XI and one of SEQ.ID.NO: 1-42 or one gene encoding PRO polypeptide or one PRO polypeptide or PRO antibody

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Inventions SEQ.ID.NO: 1-42 or one gene encoding PRO polypeptide or one PRO polypeptide or PRO antibody(if different from SEQ.ID.NO: 1-42) are not so linked as to under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: The claimed polypeptides and polynucleotides (i.e., SEQ.ID.NO 1-42) or gene encoding PRO19597, PRO83469, PROI189, PRO83470, PRO28700, PRO1246, PRO83471, PRO6244, PRO83472, PROI9600, PRO4977, PRO83473, PRO83474, PRO617, PR071057, PRO83475, PRO1065, PRO83476. PRO200, PRO1361 or PRO83477 polypeptide or PRO19597, PRO83469, PROI189, PRO83470, PRO28700, PRO1246, PRO83471, PRO6244, PRO83472, PROI9600, PRO4977, PRO83473, PRO83474, PRO617, PR071057, PRO83475, PRO1065, PRO83476, PRO200, PRO1361 or PRO83477 polypeptide or an anti-PRO19597, anti-PRO83469, anti-PRO1189, anti-PRO83470, anti-PRO28700, anti-PRO1246, anti-PRO83471, anti-PRO6244, anti-PRO83472, anti-PRO 19600, anti-PRO4977, anti-PRO83473, anti-PRO83474, anti-PRO617, anti-PRO71057, anti-PRO83475, anti-PRO 1065, anti-PRO83476, anti-PRO200, anti-PRO 1361 or anti- PRO83477 share no common special technical feature because the polynucleotides and peptides have no common structure (i.e., no common sequence) as evidenced by their sequences SEQ.ID.NO 1-68. These sequences that share no common structure as polynucleotides and polypeptides and are not linked by the same the same or a corresponding special technical feature so as to form a single general inventive concept. Therefore, where structural identity is required, such as for hybridization or expression of protein or binding of antibody, each sequence appears perform a different function in that peptides elicit an antibody response and nucleic acids encode peptides that specifically bind to an antibody. Thus they share no common structure and function so as to form a single general inventive concept under Rule 13.1. Hence, unity is lacking among groups SEQ.ID.NOS or gene encoding PRO polypeptide or PRO polypeptide or anti- PRO antibody. Applicant is required under PCT Rule 13.1 because, under PCT Rule 13.2 to elect a single disclosed SEQ.ID.NO from any group elected.

- 5. Applicant is required, in reply to this action, to elect a group and one sequence and identify the SEQ.ID.NO to which the claims shall be restricted. The reply must also identify the claims readable on the elected invention, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered non-responsive unless accompanied by an election.
- 6. Papers related to this application may be submitted to Group 1600, AU 1645 by facsimile transmission. Papers should be transmitted via the PTO Fax Center, which receives transmissions 24 hours a day and 7 days a week. The transmission of such papers by facsimile must conform to the notice

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published in the Official Gazette, 1096 OG 30, November 15, 1989. The Right Fax number is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PMR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PMR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PMR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Any inquiry concerning this communication or earlier communications from the Examiner should be directed to Padma Baskar Ph.D., whose telephone number is ((571) 272-0853. A message may be left on the Examiner's voice mail system. The Examiner can normally be reached on Monday to Friday from 6.30 a.m. to 4.00 p.m. except First Friday of each bi-week.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Jeffrey Siew can be reached on (571) 272-0787. Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (571) 272-1600

Padma Baskar Ph.D.

SUSAN UNGAR, PH.D PRIMARY EXAMINER